Effects of surface modification and SiO₂ thickness on the optical and superparamagnetic properties of the water-soluble ZnS:Mn²⁺ nanowires/Fe₃O₄ quantum dots/SiO₂ heterostructures

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In this paper, one-dimensional ZnS:Mn²⁺ nanowires (NWs)/Fe₃O₄ quantum dots (QDs)/SiO₂ heterostructures were successfully synthesized by the Stöber method to form the water-soluble fluorescent/superparamagnetic nanocomposites. The average diameter of the ZnS:Mn²⁺ NWs, Fe₃O₄ QDs and ZnS:Mn²⁺ NWs/Fe₃O₄ QDs/SiO₂ heterostructures was about 6–8 nm, 4–5 nm and 18 nm, respectively. The Fe₃O₄ QDs were covalently linked to the ZnS:Mn²⁺ NWs by the conjugation of the hydroxyl groups on the surface of the QDs and the carboxyl groups modified on the surface of the NWs. It was found that the covalent bonds between the NWs and QDs could effectively suppress the energy transfer from the ZnS:Mn²⁺ NWs to the Fe₃O₄ QDs. As the SiO₂ shell thickness increased, the fluorescence intensity reached the highest value when the hydrolysis time of tetraethyl orthosilicate was 5 hours, which was comparable to that of the ZnS:Mn²⁺ NWs. The superparamagnetic properties of the heterostructures were observed at room temperature, which decreased as the SiO₂ thickness increased.

Introduction

Next generation molecular probes combining magnetic and fluorescent properties in one entity open up broader avenues for their applications as dual-modality imaging probes in the fields of biological imaging, cell tracking, magnetic bio-separation, targeted drug delivery, etc. Among the widely studied fluorescent labeling nanomaterials, ZnS has great advantages as follows: (1) low toxicity, (2) high photostability, (3) relative lower price, (4) simple synthesis procedure. It is well known that doping Mn²⁺ ions into the ZnS lattice can give rise to the strong yellow-orange light emission located at about 583 nm (through the ⁴T₁ to ⁶A₁ internal transition) with a high quantum yield, the luminescence lifetime of which is about 1 ms. Such a strong luminescence and long lifetime allows the light penetrate into and out of the tissues and distinguish the luminescence from the background autofluorescence. These excellent optical properties make ZnS:Mn²⁺ nanocrystals potential candidates for fluorescent labeling agents.

Magnetic Fe₃O₄ nanoparticles are the most prominent class of magnetic nanoparticles (MNPs) for their biocompatibility and stability. At diameters less than 10 nm, Fe₃O₄ MNPs have unique properties such as high surface area, super-paramagnetism and low Curie temperature etc., and smaller particles may undergo rapid biodegradation when they are directly exposed to biological environments. Therefore, bifunctional optical-magnetic ZnS:Mn²⁺/Fe₃O₄ nanocomposites are highly desired in the biological field due to their appealing applications. However, fluorescence quenching always occurs due to the energy-transfer process between the ZnS:Mn²⁺ and Fe₃O₄. To solve the problem, some transition layers, e.g. silica, polymer etc. have been used in the synthesis of the core-shell nanomaterials. However, so far, the resultant nanocomposites are usually larger, and the synthetic procedure is rather complex and time-consuming. So, covalent linking between the ZnS:Mn²⁺ and Fe₃O₄ is highly desired to attenuate the energy-transfer process and increase the stability of the nanocomposites. For further bioimaging applications, it is required to transfer the hydrophobic nanocomposites to water by surface modification, such as polymer, silica, or carbon coating techniques. Among them, silica coating has attracted a great deal of attention because of its excellent chemical stability, biocompatibility and facile further conjugation with various functional groups. To date, only a few
methods have been reported for the preparation of water-soluble silica-coated nanocomposites with a size below 20 nm.

In this paper, we firstly designed a facile approach for fabricating bi-functional optical-magnetic ZnS:Mn\(^{2+}\) nanowires (NWs) decorated by Fe\(_3\)O\(_4\) quantum dots (QDs) with different thickness of SiO\(_2\) without using other components as the transition layer. The thioglycic acid modified ZnS:Mn\(^{2+}\) NWs were conjugated with the citric acid modified Fe\(_3\)O\(_4\) QDs to form stable nanocomposites through the reaction between the hydroxyl groups on the surface of the QDs and the carboxyl groups modified on the surface of the NWs. The effects of the surface modification and SiO\(_2\) thickness on the optical and superparamagnetic properties of the nanocomposites were investigated.

### Experimental section

#### Materials

Zinc nitrate, manganese nitrate, thiourea, ethylenediamine (EN), thioglycic acid (MPA), ferric chloride (FeCl\(_3\)-6H\(_2\)O), ferrous chloride (FeCl\(_2\)-4H\(_2\)O), PEG-4000, ethylene glycol, concentrated ammonia aqueous solution (25%), citric acid, tetraethyl orthosilicate (TEOS) and ethanol were all analytical grade (Shanghai Chemical Reagents Co.), and used without further purification.

#### Preparation and chemical modification of ZnS:Mn\(^{2+}\) (1%) NWs

In a typical process, 0.99 mmol of zinc nitrate and 0.01 mmol of manganese nitrate were dissolved in 16 ml EN and water (1:1 in volume ratio). After stirring for 1 hour, 3 mmol of thiourea was added to the resulting complex. After stirring for 2 hours, the colloid solution was transferred into a 20 ml Teflon-lined autoclave and kept at 180 °C for 12 hours. After the reaction, the autoclave was taken out and cooled down to room temperature. The product was washed with ethanol and deionized water for several times and separated by centrifugation, and then dried at 80 °C for 1 hour to yield a white powder.

ZnS:Mn\(^{2+}\) (1%) NWs was modified with MPA as follows: 9.7 mg of ZnS:Mn\(^{2+}\) NWs were dispersed ultrasonically in deionized water (100 ml), and then 75 μl of MPA was added to the above solution under magnetic stirring for 1 hour to form the ZnS:Mn\(^{2+}\)(–SH) water solution.

#### Preparation and chemical modification of Fe\(_3\)O\(_4\) QDs

Fe\(_3\)O\(_4\) QDs were prepared through an improved chemical coprecipitation method. FeCl\(_3\)-6H\(_2\)O (6 mmol), FeCl\(_2\)-4H\(_2\)O (10 mmol) and PEG-4000 (10 g) were dissolved in absolute ethylene glycol (250 ml). After stirring for 30 min at room temperature, 100 ml of concentrated ammonia aqueous solution (25%) was added rapidly to the resulting complex. Then the reaction mixture was heated at 110 °C for 2 hours under mechanical stirring, producing a black and homogeneous suspension. After cooling to room temperature, the precipitate was separated with a permanent magnet and washed with deionized water and ethanol for several times until the pH decreased to 7.0, followed by drying in a vacuum oven at 40 °C for 48 hours.

Fe\(_3\)O\(_4\) QDs was modified with citric acid as follows: 0.023 g of Fe\(_3\)O\(_4\) QDs and 0.023 g of citric acid were dispersed in 250 ml of deionized water under mechanical stirring for 1 hour to form the Fe\(_3\)O\(_4\)(–COOH) water solution.

#### Preparation of ZnS:Mn\(^{2+}\)(–SH) NWs/Fe\(_3\)O\(_4\)(–COOH) QDs/SiO\(_2\) core–shell nanostructures

100 ml of ZnS:Mn\(^{2+}\)(–SH) water solution and 250 ml of Fe\(_3\)O\(_4\)(–COOH) water solution were mixed in a three-necked flask under mechanical stirring for 1 hour. A mixed solution including 50 ml of absolute alcohol, 1 ml of distilled water, 1.7 ml of aqueous ammonia, and 200 μl of TEOS was injected into the above solution. The mixture was continuously stirred for 5 hours at room temperature. The product was washed with ethanol several times and separated by centrifugation, and then dried at 60 °C for 3 hours to obtain a powder. To investigate the effects of the SiO\(_2\) thickness on the optical and magnetic properties of the samples, a set of samples were synthesized at different time periods (t = 1, 3, 7 hours). The same process was carried out except for using the ZnS:Mn\(^{2+}\) and Fe\(_3\)O\(_4\) to displace the ZnS:Mn\(^{2+}\)(–SH) and Fe\(_3\)O\(_4\)(–COOH) for comparison.

#### Characterization of products

X-ray diffraction (XRD) patterns were collected on a MAC Science MXP-18 X-ray diffractometer using a Cu target radiation source. Transmission electron micrographs (TEM) and high-resolution transmission electron microscopy (HRTEM) images were taken on a JEM-2100 electron microscope. The specimens were prepared by depositing a drop of the dilute solution of the sample in ethanol on a carbon-coated copper grid and drying at room temperature. EDAX microanalysis was performed at the SEM magnification. Fourier transform infrared (FT-IR) spectra were recorded on a Bruker Vertex 70 spectrophotometer in KBr pellets. Photoluminescence (PL) measurements were carried out at room temperature, using 325 nm as the excitation wavelength and a He–Cd laser as the source of excitation. Magnetic hysteresis loop was measured by a Lake Shore 7407 vibrating sample magnetometer (VSM) with the maximum field of 6 kOe. The samples used for the XRD, EDAX, FT-IR, PL and VSM measurements were solid powders.

The quantum yield (QY) of the sample was measured relative to the organic dye Rhodamine B (5.0 μg mL\(^{-1}\)) in ethanol solution (QY = 95%), which can be determined from the following equation:

\[
Q_Y = Q_Y^{(s)} \left( \frac{E_s}{E_x} \right) \left( \frac{A_x}{A_s} \right) \left( \frac{n_x}{n_s} \right)^2
\]

In the equation, \(E_s\) (sample) and \(E_x\) (standard) are the integrated emission peak areas, \(A_x\) (sample) and \(A_s\) (standard) are the absorbance at the excitation wavelength, and \(n_x\) (sample) and \(n_s\) (standard) are the refractive indices of the solvents: \(n_{ethanol}\) is 1.359 and \(n_{water}\) is 1.333 at room temperature. UV-Vis absorption spectrum was measured on a UV-3101PC UV spectrometer. PL spectrum was taken using a fluorescence spectrophotometer (Perkin-Elmer, LS55) at room temperature. The specimen for the
measurement was dispersed in water and placed in a 1 cm quartz cell, and water served as the reference. In order to reduce the measurement error, multiple concentrations were examined for the sample, and the optical densities at the excitation wavelengths of both Rhodamine B and the sample solution were set to be always below 0.08 to minimize the reabsorption effect.

Results and discussion

Fig. 1 shows the XRD patterns of the ZnS:Mn$^{2+}$ NWs (black line); Fe$_3$O$_4$ QDs (blue line); ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites (red line).

The detailed microstructures of the prepared samples are characterized using transmission electron microscopy. Fig. 2a and b display the TEM, EDAX and HRTEM images of the ZnS:Mn$^{2+}$ NWs. The HRTEM image (Fig. 2b) was taken from the area marked by the red rectangle in Fig. 2a. It can be seen that the ZnS:Mn$^{2+}$ NWs are smooth and the diameter is about 6–8 nm. The growth direction of the ZnS:Mn$^{2+}$ NWs is perpendicular to the lattice fringes, and the $d$ spacing of the (002) plane is about 0.31 nm. The EDAX image of the ZnS:Mn$^{2+}$ NWs (the inset figure in Fig. 2a) shows that the sample contains Zn, S and Mn elements, 0.56 at% Mn can be detected for the ZnS:Mn$^{2+}$ (1%) NWs. Fig. 2c and d show the TEM, size distribution histogram and HRTEM images of the Fe$_3$O$_4$ QDs. It can be seen that these QDs are well crystallized and the average diameter of the QDs is about 4–5 nm. The lattice fringes with the $d$ spacing of 0.25 nm can be assigned to the (311) plane of the cubic Fe$_3$O$_4$ QDs, which is consistent with the XRD results.

Fig. 3 displays the TEM and HRTEM images of the ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites. The TEM image (Fig. 3a) shows an apparent contrast between the inner core and the outer shell, the QDs assemble uniformly around the surface of the NWs, which suggests the existence of the core–shell structure. The average diameter of the ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites is about 18 nm, which is not obviously enlarged, revealing that the SiO$_2$ layer is thin. The different lattice fringes between the Fe$_3$O$_4$ QDs and ZnS:Mn$^{2+}$ NWs can be clearly identified in the HRTEM image (Fig. 3b) taken from the area marked by the red rectangle in Fig. 3a, which provides strong evidence for the composite nature of the sample.

Fig. 4 shows the proposed formation mechanism of the ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites. For the ZnS:Mn$^{2+}$ NWs modified by MPA, thiol groups (–SH) can form stable bonds with the metals (Zn$^{2+}$) with the carboxyl groups (–COOH) protruding outwards from the surface of the NWs. Fe$_3$O$_4$ QDs are functionalized with citric acid, the surface of which are modified by carboxyl groups (–COOH) and hydroxyl groups (–OH). After mixing the ZnS: Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs together, the QDs can be covalently bonded to the NWs by the –COO groups in order to avoid the charge transfer from the ZnS:Mn$^{2+}$ NWs to the Fe$_3$O$_4$ QDs, thereby retaining both the fluorescence and magnetic properties. Subsequently, silica is coated on the surface of Fe$_3$O$_4$ QDs to form the ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites through the hydrolysis of TEOS according to the Stöber method. The SiO$_2$ shell can effectively prevent the aggregation and chemical degradation of the ZnS:Mn$^{2+}$/Fe$_3$O$_4$ nanocomposites in a harsh liquid environment.

FT-IR spectra were recorded and used to identify the changes of the functional groups on the surface of the nanocomposites. For the Fe$_3$O$_4$ QDs (red line in Fig. 5), the vibration of the Fe–O bonds centered at 585.9 cm$^{-1}$ is the characteristic absorption of the Fe$_3$O$_4$. For the ZnS:Mn$^{2+}$/Fe$_3$O$_4$/SiO$_2$ nanocomposites (green line in Fig. 5), the strong peak at 1095.7 cm$^{-1}$ can be assigned to the vibration of the Si-
O bonds, demonstrating the existence of the SiO2 components. For the ZnS:Mn2+/Fe3O4/SiO2 nanocomposites (blue line in Fig. 5), the two peaks at 1406 and 1576 cm⁻¹ correspond to the stretching vibration of the carboxyl, which indicates that the citric acid have been successfully modified on the surface of the Fe3O4 QDs. For three samples, a strong peak at 3411 cm⁻¹ can be observed, which is ascribed to the stretching vibration of the O–H bonds in water. Moreover, the intensity of the Fe–O bonds for the ZnS:Mn2+/Fe3O4/SiO2 and ZnS:Mn2+(SH)/Fe3O4(COOH)/SiO2 nanocomposites is much lower than that for the Fe3O4 QDs, which further proves that the Fe3O4 QDs have been coated by citric acid and SiO2.

Surface modification always plays a critical role in determining the optical properties of the nanocrystals. Fig. 6a shows the room temperature PL spectra of the ZnS:Mn2+ NWs, ZnS:Mn2+/Fe3O4/SiO2, ZnS:Mn2+(SH)/Fe3O4(SiO2, and ZnS:Mn2+(SH)/Fe3O4(COOH)/SiO2 nanocomposites. The PL spectrum of the ZnS:Mn2+ NWs (black line in Fig. 6a) shows a strong yellow-orange emission peak at 583 nm (corresponding to the Mn2⁺ 4T1g→4A1g transition), indicating...
that the Mn$^{2+}$ ions have been successfully incorporated into the ZnS lattice. After embedding the ZnS:Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs together within the SiO$_2$ matrix, the intensity of the yellow-orange emission (red line in Fig. 6a) is approximately three times lower than that of the unpassivated counterpart (black line in Fig. 6a). Since that the interaction between the ZnS:Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs would lead to the energy transfer from the NWs to the QDs, hence reduce the PL intensity. After coating –SH groups on the surface of the NWs, the intensity of the ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$/SiO$_2$ nanocomposites is further decreased (green line in Fig. 6a). Zheng et al. have pointed out that the thiol ligands acted as the PL quencher for the ZnS:Mn$^{2+}$ QDs due to the lower energy levels of the valence band ($\approx 1.78$ eV) compared to the highest occupied molecular orbital levels of the thiol ligands ($\approx 1.0$ eV). It is worth noting that the PL intensity of the as-synthesized ZnS:Mn$^{2+}$(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites (blue line in Fig. 6a) is comparable to that of the ZnS:Mn$^{2+}$ NWs (black line in Fig. 6a) after modifying the Fe$_3$O$_4$ QDs with citric acid. The PL enhancement can be attributed to the following factors: firstly, the dominant mechanism could be the changed surface structure and the effective surface passivation induced by the ligand exchange. So there would be more energy transfer from the hosts to the d-orbital of the Mn$^{2+}$ ions than that to the nonradiative recombination centers on the surface.
of the ZnS:Mn$^{2+}$ NWs; secondly, the ligands (–OH, –COOH, –COO) between the NWs and QDs can serve as an extra barrier for confining the electrons to the interior of the NWs, which could increase the possibility of the electrons to fall into the d-orbital, and subsequently enhance the dopant emission.$^{36}$ Moreover, the QY of the ZnS:Mn$^{2+}$/(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites is calculated to be 20±1%, indicating that the nanocomposites possess high quality luminescent properties. In addition, the PL spectrum of the ZnS:Mn$^{2+}$/(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites becomes asymmetric due to the joints between the NWs and QDs, which is consistent with the results of CdS/Fe$_3$O$_4$ heterostructures.$^{37}$ It can be deconvoluted into six Gaussian peaks centered at 401, 420, 445, 498, 538, 574 nm, respectively. According to the energy diagram of the defects distributed in the ZnS, the emission peaks can be attributed to the following origins: 401 nm to the sulfur vacancy and interstitial sulfur lattice defects,$^{38}$ 420 and 445 nm to the sulfur vacancy and surface states,$^{39,40}$ 498 nm to the self-activated defect centers formed by the zinc vacancy inside the lattice,$^{41}$ 538 nm to the sulfur species on the surface of the ZnS NWs,$^{42}$ and 574 nm to the Mn$^{2+}$$^4T_1$$^6A_1$ transition.$^{32}$ It is noticeable that the yellow-orange emission shifts from 585 nm to 574 nm after modifying the Fe$_3$O$_4$ QDs with citric acid, i.e. blue shift occurs. The interactive straining caused by the large variation in lattice parameters between the core and shell may lead to the blue shift of the yellow-orange emission.

To investigate the effect of the SiO$_2$ shell thickness on the optical properties of the ZnS:Mn$^{2+}$/(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites, the PL spectra of the samples grown at different times are shown in Fig. 6b. For the yellow-orange emission, the PL intensity keeps increasing until $t = 5$ h and then decreases rapidly when the time continually increases to 7 h. The PL intensity of the emission is mainly determined by the degree of surface passivation. At the early stage of the hydrolysis of TEOS, only parts of ZnS:Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs were encapsulated by the silica shell, while the rest of the ZnS:Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs still remained in the solution, which can be removed in the experiment. So, the surface of the ZnS:Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs can be passivated fully by the Fe$_3$O$_4$ QDs and SiO$_2$, respectively. With the increase in the hydrolysis time of TEOS, the silica shell became thick and more ZnS:Mn$^{2+}$ NWs and Fe$_3$O$_4$ QDs would be encapsulated. But the thick Fe$_3$O$_4$ and SiO$_2$ coating would induce strain at the interface caused by the lattice mismatch between one another. It would result in new traps for carriers, which are thought to be the main reason for the decrease of the yellow-orange emission after $t = 5$ h.

The magnetic properties of the Fe$_3$O$_4$ QDs, the ZnS:Mn$^{2+}$/(SH)/Fe$_3$O$_4$(COOH)/SiO$_2$ nanocomposites with different SiO$_2$ thickness, ZnS:Mn$^{2+}$/(SH)/Fe$_3$O$_4$/SiO$_2$ and ZnS:Mn$^{2+}$/Fe$_3$O$_4$/SiO$_2$ nanocomposites measured at room temperature are shown in Fig. 7. All of the samples exhibit superparamagnetic properties, where negligible coercivity and remanence in the magnetization curves are observed and the magnetic moments are only induced in the presence of a magnetic field. The magnetic saturation ($M_s$) value of the Fe$_3$O$_4$ QDs is 20 emu g$^{-1}$ (Fig. 7a), which is lower than that of the bulk Fe$_3$O$_4$ (92 emu g$^{-1}$).$^{43}$ Since the saturation magnetization as well as the magnetic moment of the particle is proportional to its size,$^{44}$ hence the as-synthesized Fe$_3$O$_4$ QDs exhibit the smaller $M_s$. However, the $M_s$ value is reduced largely in the presence of ZnS:Mn$^{2+}$ NWs and amorphous SiO$_2$ compared with the Fe$_3$O$_4$ QDs (Fig. 7b and c), which is
consistent with the literature.\textsuperscript{45} For the ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}(COOH)/SiO\textsubscript{2} nanocomposites (Fig. 7b), the $M_s$ is decreased as the thickness of the SiO\textsubscript{2} increased, which is mainly due to the lower mass content of Fe\textsubscript{3}O\textsubscript{4} in the nanocomposites as the mass of the SiO\textsubscript{2} increased.\textsuperscript{45} It is noted that the surface modification also play an important role in affecting the magnetic properties, which can be seen in Fig. 7c. The decreased $M_s$ value for the ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}/SiO\textsubscript{2} and ZnS:Mn\textsuperscript{2+}/Fe\textsubscript{3}O\textsubscript{4}/SiO\textsubscript{2} nanocomposites compared with the ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}(COOH)/SiO\textsubscript{2} nanocomposites can be attributed to the surface spin disorder due to capping.\textsuperscript{13} In addition, surface modification can also make the NWs load more QDs on its surface by the stable covalent bonds.

Fig. 8 shows the photographic images of the ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}(COOH)/SiO\textsubscript{2} nanocomposites under white light and UV illumination ($\lambda = 365$ nm) with and without an external magnetic field. It can be seen that the solution of the as-synthesized nanocomposites is orange and the nanocomposites are well dispersed in the aqueous solution under white light (Fig. 8a and c) and UV irradiation (Fig. 8d and f). When a permanent magnet is placed near the solution (Fig. 8b and e), the nanocomposites are attracted and accumulated toward the magnet, eventually leaving a clear solution behind, indicating that the magnetic separation occurs. From the optical and magnetic properties of the nanocomposites, it can be concluded that the ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}(COOH)/SiO\textsubscript{2} nanocomposites show the strong fluorescent emission and desired superparamagnetic properties, which enable them to serve as multifunctional nanocomposites with promising applications in biological imaging, magnetic guiding and separation.

Conclusions

In this paper, the water-soluble ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}(COOH)/SiO\textsubscript{2} nanocomposites were fabricated successfully by assembling the citric acid capped Fe\textsubscript{3}O\textsubscript{4} QDs on MPA-coated ZnS:Mn\textsuperscript{2+} NWs by the Stöber method. The effects of the surface modification and SiO\textsubscript{2} thickness on its optical and magnetic properties were studied. It was found that the citric acid coating on the surface of the Fe\textsubscript{3}O\textsubscript{4} QDs could effectively suppress the interaction between the ZnS:Mn\textsuperscript{2+} NWs and Fe\textsubscript{3}O\textsubscript{4} QDs. The presented water-soluble core–shell nanocomposites offer the excellent yellow-orange emission of the ZnS:Mn\textsuperscript{2+} NWs and retain the superparamagnetic properties of the Fe\textsubscript{3}O\textsubscript{4} QDs. Therefore, the fluorescent, superparamagnetic and water-soluble properties of the ZnS:Mn\textsuperscript{2+}(SH)/Fe\textsubscript{3}O\textsubscript{4}(COOH)/SiO\textsubscript{2} nanocomposites would allow them to find applications in biolabeling, bioseparation and diagnostic analysis.

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